

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances Application: United States Pharmacopeial

Convention

[Docket No. DEA-392]

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER]. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

ADDRESS: Written comments should be sent to: Drug Enforcement Administration,
Attention: DEA Federal Register Representative/ODW, 8701 Morrissette Drive,
Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement
Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia
22152. All requests for hearing should also be sent to: (1) Drug Enforcement
Administration, Attn: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia
22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register
Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION:

The Attorney General has delegated her authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control ("Deputy Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on February 26, 2016, United States Pharmacopeial Convention, 12601 Twinbrook Parkway, Rockville, Maryland 20852 applied to be registered as an importer of the following basic classes of controlled substances:

| Controlled Substance | Schedule |
|--|-----------------|
| Cathinone (1235) | I |
| Methaqualone (2565) | I |
| Lysergic acid diethylamide (7315) | I |
| Marihuana (7360) | I |
| Tetrahydrocannabinols (7370) | I |
| 4-Methyl-2,5-dimethoxyamphetamine (7395) | I |
| 3,4-Methylenedioxyamphetamine (7400) | I |
| Codeine-N-oxide (9053) | I |
| Difenoxin (9168) | I |
| Heroin (9200) | I |
| Morphine-N-oxide (9307) | I |
| Norlevorphanol (9634) | I |
| Methamphetamine (1105) | II |
| Phenmetrazine (1631) | II |
| Methylphenidate (1724) | II |
| Amobarbital (2125) | II |
| Pentobarbital (2270) | II |

| Controlled Substance | Schedule |
|--|-----------------|
| Secobarbital (2315) | II |
| Glutethimide (2550) | II |
| Phencyclidine (7471) | II |
| 4-Anilino-N-phenethyl-4-piperidine (ANPP) (8333) | II |
| Phenylacetone (8501) | II |
| Alphaprodine (9010) | II |
| Anileridine (9020) | II |
| Cocaine (9041) | II |
| Dihydrocodeine (9120) | II |
| Diphenoxylate (9170) | II |
| Levomethorphan (9210) | II |
| Levorphanol (9220) | II |
| Meperidine (9230) | II |
| Dextropropoxyphene, bulk (non-dosage forms) (9273) | II |
| Thebaine (9333) | II |
| Noroxymorphone (9668) | II |
| Alfentanil (9737) | II |
| Sufentanil (9740) | II |

The company plans to import the listed controlled substances in bulk powder form from foreign sources for the manufacture of analytical reference standards for sale to their customers.

The company plans to import analytical reference standards for distribution to its customers for research and analytical purposes. Placement of these drug codes onto the company's registration does not translate into automatic approval of subsequent permit applications to import controlled substances. Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under to 21 U.S.C. 952(a)(2). Authorization will not extend to the import of FDA approved or non-approved finished dosage forms for commercial sale.

Dated: September 7, 2016

Louis J. Milione,

Deputy Assistant Administrator.

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